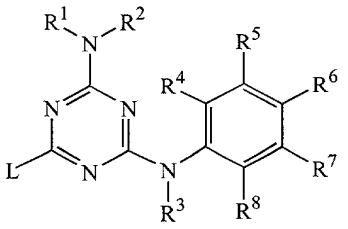
ABSTRACT

SUBSTITUTED DIAMINO-1,3,5-TRIAZINE DERIVATIVES

This invention concerns the compounds of formula



(I)

the pharmaceutically acceptable acid addition salts and the stereochemically isomeric forms thereof, wherein R^1 and R^2 are each independently selected from hydrogen; hydroxy; amino; optionally substituted C_{1-6}alkyl; C_{1-6}alkyloxy; C_{1-6}alkylcarbonyl; C_{1-6}alkyloxycarbonyl; Ar^1; mono- or di(C_{1-6}alkyl)amino; mono- or di(C_{1-6}alkyl)-aminocarbonyl; dihydro-2(3*H*)-furanone; or R^1 and R^2 taken together may form pyrrolidinyl, piperidinyl, morpholinyl, azido or mono- or di(C_{1-6}alkyl)amino-C_{1-4}alkylidene; R^3 is hydrogen, Ar^1, C_{1-6}alkylcarbonyl, C_{1-6}alkyl, C_{1-6}alkyloxy-carbonyl, C_{1-6}alkyl substituted with C_{1-6}alkyloxycarbonyl; and R^4, R^5, R^6, R^7 and R^8 are each independently selected from hydrogen, halo, C_{1-6}alkyl, C_{1-6}alkyloxy, cyano, aminocarbonyl, nitro, amino, trihalomethyl or trihalomethyloxy; L is optionally substituted C_{1-10}alkyl; C_{3-10}alkenyl; C_{3-10}alkynyl; C_{3-7}cycloalkyl; Ar^1 is optionally substituted phenyl; for the manufacture of a medicine for the treatment of subjects suffering from HIV (Human Immunodeficiency Virus) infection. It further relates to new compounds being a subgroup of the compounds of formula (I), their preparation and compositions comprising them.